

CAB17454

Product Information

Product SKU:	CAB17454	Gene ID:	4715		Size:	20uL, 100uL	
Clone No:	-	Host Species:	Rabbit		Reactivity :	Human, Mouse, Rat	
Additional Information							
Observed MW	22kDa		Conjugato:	Unconiugated	4		

Observed MW:	22kDa	Conjugate:	Unconjugated
Calculated MW:	22kDa	lsotype:	lgG

Immunogen Information

Background	The protein encoded by this gene is a subunit of the mitochondrial oxidative phosphorylation complex
	I (nicotinamide adenine dinucleotide: ubiquinone oxidoreductase). Complex I is localized to the inner
	mitochondrial membrane and functions to dehydrogenate nicotinamide adenine dinucleotide and to
	shuttle electrons to coenzyme Q. Complex I deficiency is the most common defect found in oxidative
	phosphorylation disorders and results in a range of conditions, including lethal neonatal disease,
	hypertrophic cardiomyopathy, liver disease, and adult-onset neurodegenerative disorders. Pseudogenes
	of this gene are found on chromosomes five, seven and eight. Alternative splicing results in multiple
	transcript variants.
Recommended Dilution:	WB,1:500 - 1:1000 IHC-P,1:50 - 1:200
Synonyms:	B22; LYRM3; CI-B22; UQOR22; MC1DN24; NDUFB9
Purifcation Method:	Affinity purification
Immunogen:	Recombinant fusion protein containing a sequence corresponding to amino acids 80-170 of human
	NDUFB9 (NP_004996.1).
Storage:	Store at -20°C. Avoid freeze / thaw cycles.Buffer: PBS with 0.01% thimerosal,50% glycerol,pH7.3.